Join Regulatory Affairs

Are you one of tomorrow’s leaders within drug development?
Use your scientific life science degree to kick-start your career with us.

What is it about?
In Novo Nordisk, Regulatory Affairs secures the approval of new multi-billion dollar products and drug indications, making it possible for the company to bring innovative products to patients. We stand at the centre of project planning and execution from the earliest stages of drug development all the way through life cycle management of marketed products. The Regulatory Affairs Graduate Programme has been designed to develop top talents in becoming future leaders within this area.

What is in it for me?
Novo Nordisk is a successful global pharmaceutical company and the world leader in diabetes care. Working at Novo Nordisk will give you the opportunity to make a significant difference to millions of patients around the world, while also delivering exciting results for the business.

During the Regulatory Affairs Graduate Programme, you will:
• Receive formal training and gain knowledge on the key role Regulatory Affairs plays in drug development and invaluable international experience
• Develop your core skills within areas such as developing and submitting marketing applications for new products, product changes and clinical trials while working closely with other areas in Novo Nordisk
• Work with senior managers and build an international network of world-class colleagues

We’ve designed the programme so that, following its successful completion, you will be equipped with the necessary skills and experience to become an integral part of our company – typically with a permanent position in Denmark or elsewhere in our global organisation.

2-year programme: three 8-month rotations
During the 2-year programme you will undertake two rotations in our headquarters in Denmark and one rotation in one of our global affiliates. You will work as a full member of a team, with responsibilities for one or more projects, and play an instrumental role in ensuring that the Novo Nordisk Regulatory Affairs organisation realizes its vision of achieving and sustaining best in class approvals to make a difference for patients.

< 2 months in:

> 4 months in:

Elisabeth Buhl Thubron, RA Graduate

Am I qualified?
To apply for the Regulatory Affairs graduate programme, you must have:
• A Master’s or PhD degree from 2018 or 2019 in a natural science
• A minimum of 6 months of international experience from working, studying or voluntary work
• Relevant work or voluntary experience (preferably no more than one year of professional work experience since finishing your studies)
• Above average academic achievements
• Professional fluency in English

Apply now
If you meet all the criteria listed in ‘Am I qualified?’, then apply before January 2019. In addition to completing the online application, you must provide a 1-minute video of yourself explaining why you are the ideal candidate for the Regulatory Affairs graduate programme.

Want to be a Regulatory Affairs Graduate in 2019?
Apply before January 2019